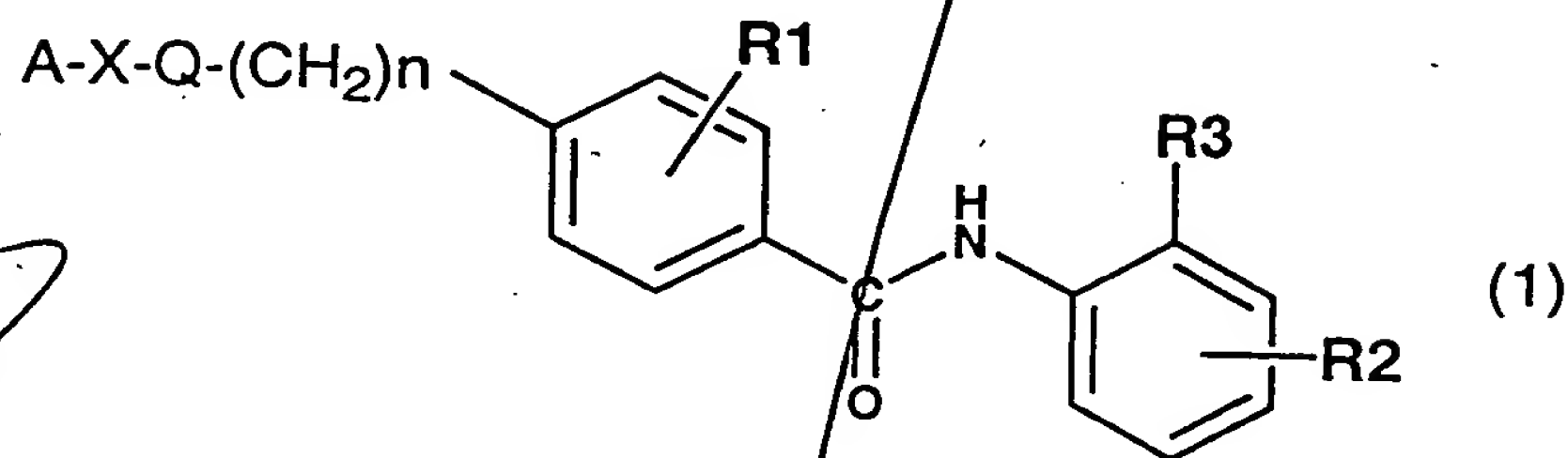


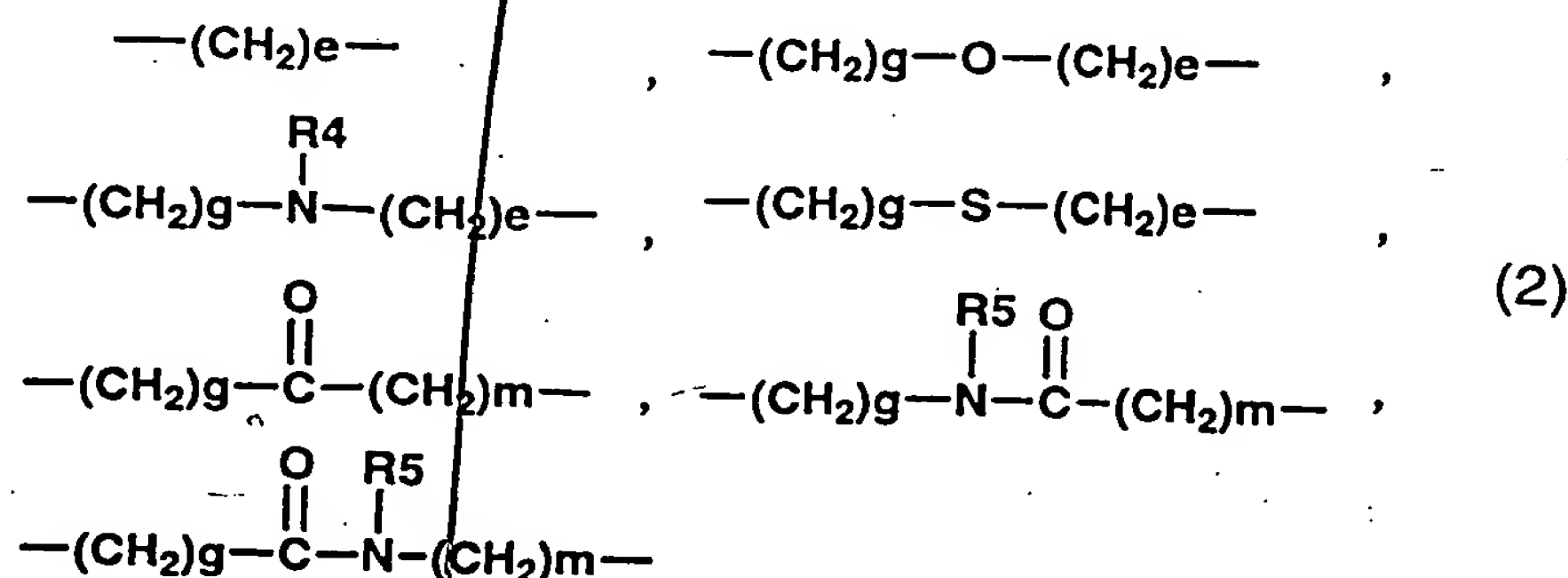
WHAT IS CLAIMED IS:

1. A compound represented by formula (1):

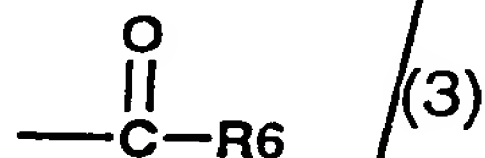


wherein A is an optionally substituted a phenyl or heterocyclic group which has 1 to 4 substituents selected from the group consisting of a halogen atom, a hydroxyl group, an amino group, a nitro group, a cyano group, an alkyl group having 1 to 4 carbons, an alkoxy group having 1 to 4 carbons, an aminoalkyl group having 1 to 4 carbons, an alkylamino group having 1 to 4 carbons, an acyl group having 1 to 4 carbons, an acylamino group having 1 to 4 carbons, an alkylthio group having 1 to 4 carbons, a perfluoroalkyl group having 1 to 4 carbons, a perfluoroalkyloxy group having 1 to 4 carbons, a carboxyl group, an alkoxycarbonyl group having 1 to 4 carbons, a phenyl group and a heterocyclic group;

X is a bond or a moiety having a structure selected from those illustrated in formula (2):



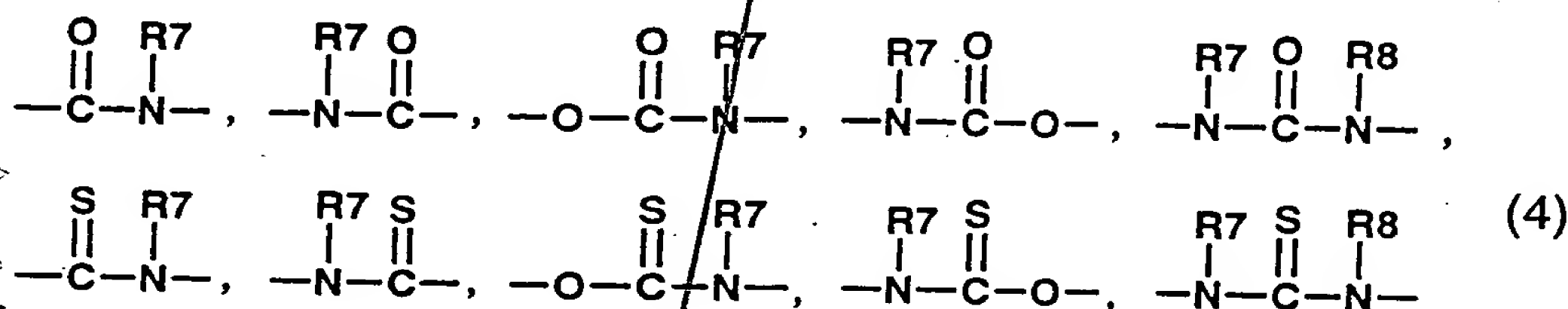
wherein e is an integer of 1 to 4; g and m are independently an integer of 0 to 4; R⁴ is a hydrogen atom or an optionally substituted alkyl group having 1 to 4 carbons, or the acyl group represented by formula (3)



wherein R⁶ is an optionally substituted alkyl group having 1 to 4 carbons, a perfluoroalkyl group having 1 to 4 carbons, a phenyl group or a heterocyclic group; R⁵ is a hydrogen atom or an optionally substituted alkyl group having 1 to 4 carbons;

n is an integer of 0 to 4, provided that when x is a bond, n is not zero;

Q is a moiety having a structure selected from those illustrated in formula (4)



wherein R⁷ and R⁸ are independently a hydrogen atom or an optionally substituted alkyl group having 1 to 4 carbons;

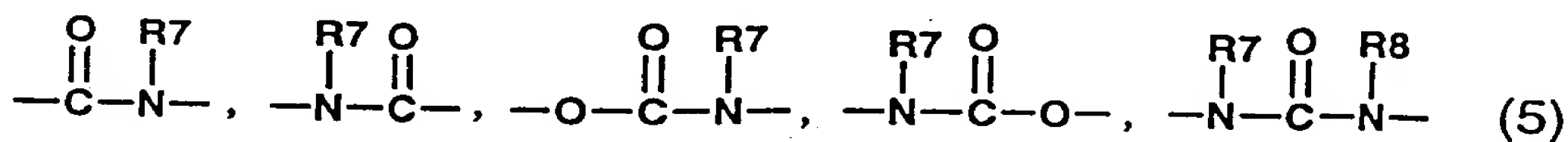
R¹ and R² are independently a hydrogen atom, a halogen atom, a hydroxyl group, an amino group, an alkyl group having 1 to 4 carbons, an alkoxy group having 1 to 4 carbons, an aminoalkyl group having 1 to 4 carbons, an alkylamino group having 1 to 4 carbons, an acyl group having 1 to 4 carbons, an acylamino group having 1 to

4 carbons, an alkylthio group having 1 to 4 carbons, a perfluoroalkyl group having 1 to 4 carbons, a perfluoroalkyloxy group having 1 to 4 carbons, a carboxyl group or an alkoxycarbonyl group having 1 to 4 carbons;

R^3 is a hydroxyl or amino group or a pharmaceutically acceptable salt thereof.

2. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 1, wherein n is an integer of 1 to 4.

3. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 2, wherein Q is selected from the structures illustrated in formula (5):



wherein R^7 and R^8 are as defined above.

4. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 3, wherein A is an optionally substituted hetero ring.

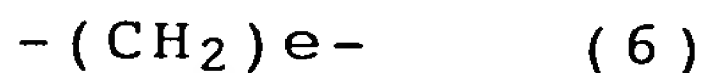
5. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 4, wherein A is an optionally substituted pyridyl group.

6. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 4, wherein X is a direct bond.

7. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 6, wherein R^1 and R^2 are a hydrogen atom.

8. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 7, wherein R^3 is an amino group.

9. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 5, wherein X is the structure represented by formula (6):

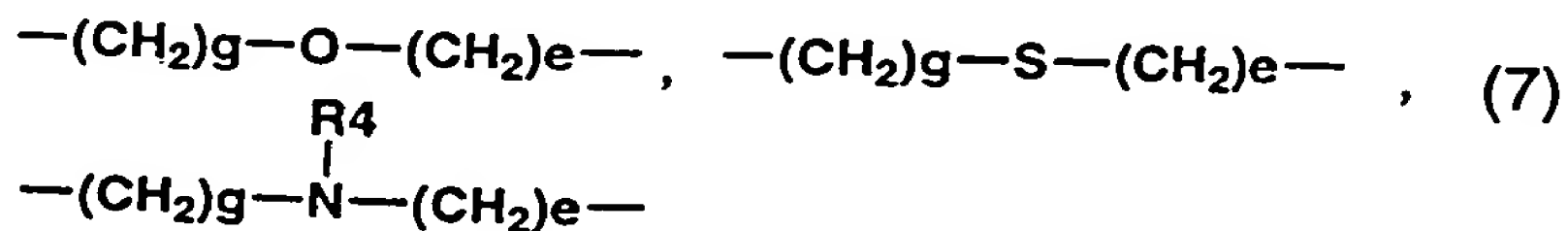


an integer of 1 to 4
~~wherein e is as defined above.~~

10. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 9, wherein n is 1; and R^1 and R^2 are a hydrogen atom.

11. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 10, wherein R^3 is an amino group.

12. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 5, wherein X is selected from the structures illustrated in formula (7):

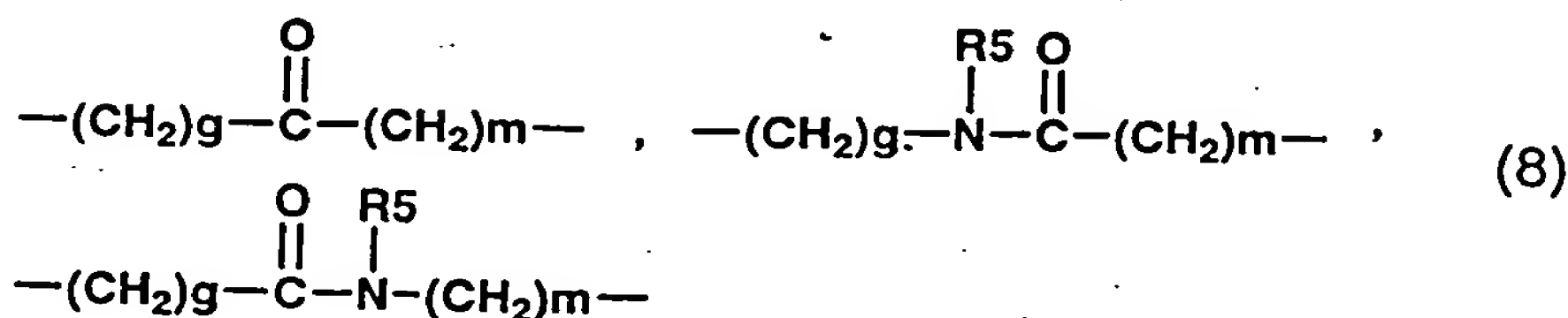


~~wherein e, g and R^4 are as defined above.~~

13. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 12, wherein n is 1; and R^1 and R^2 are a hydrogen atom.

14. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 13, wherein R^3 is an amino group.

15. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 5, wherein X is selected from the structures illustrated in formula (8):



wherein g, m and R⁵ are as defined above.

16. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 15, wherein n is 1; and R¹ and R² are a hydrogen atom.

17. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 16, wherein R³ is an amino group.

18. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 1, wherein n is zero.

19. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 18, wherein Q is selected from the structures illustrated in formula (5).

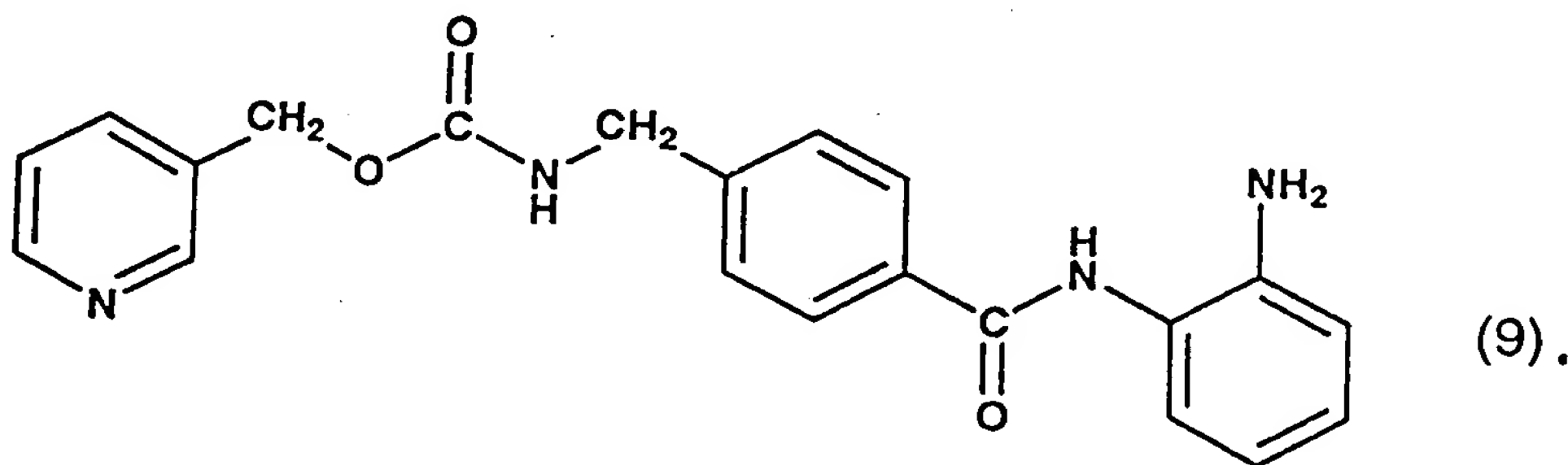
20. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 19, wherein A is an optionally substituted hetero ring.

21. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 20, wherein A is an optionally substituted pyridyl group.

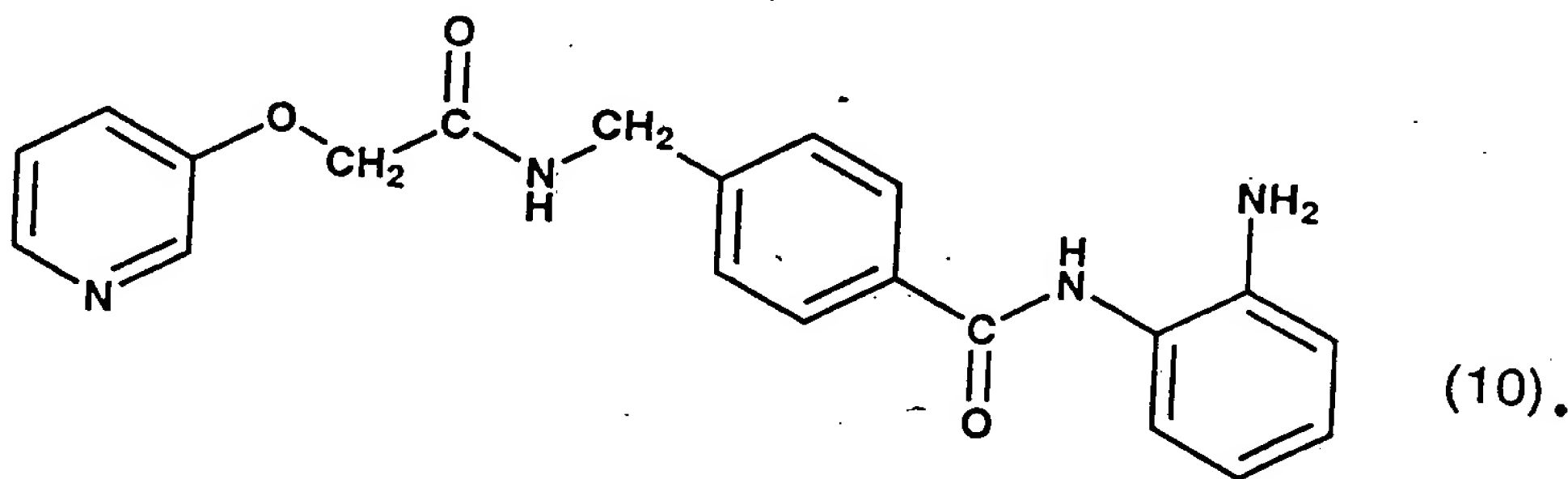
22. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 21, wherein R^1 and R^2 are a hydrogen atom.

23. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 22, wherein R^3 is an amino group.

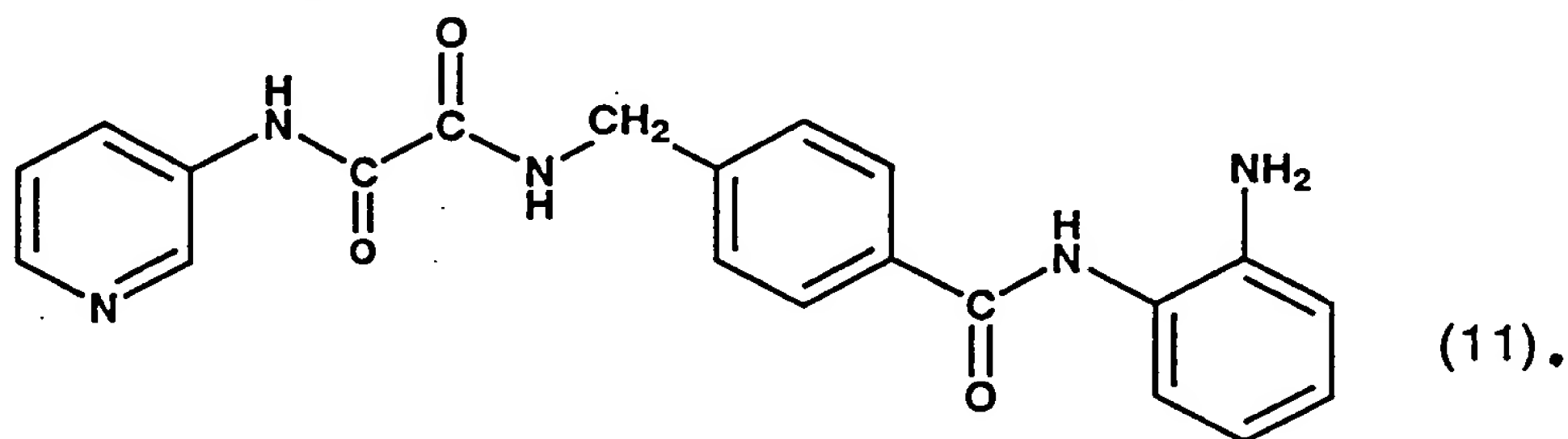
24. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 1 represented by formula (9).



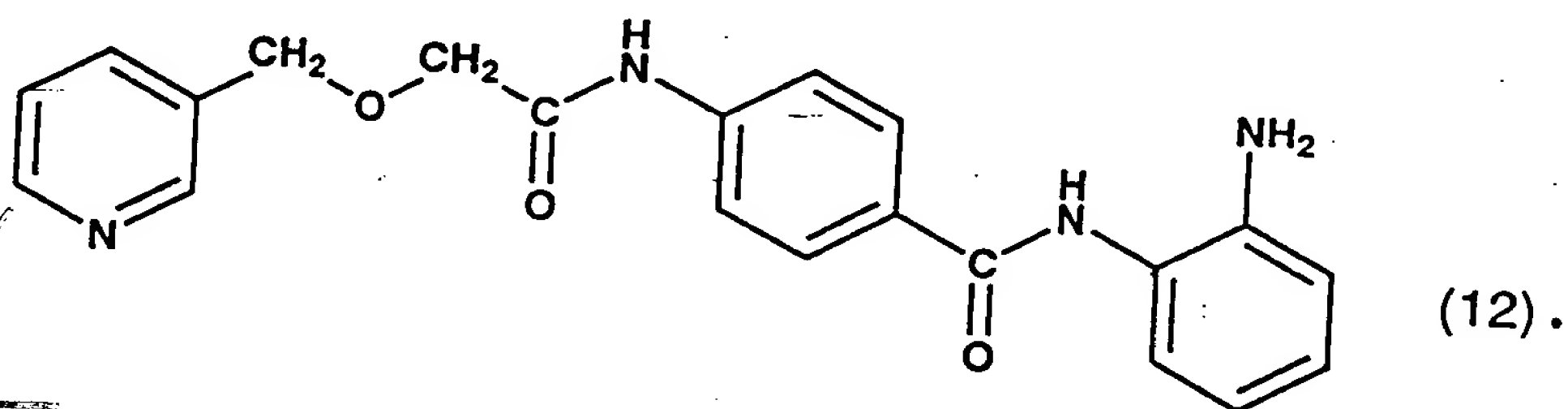
25. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 1 represented by formula (10).



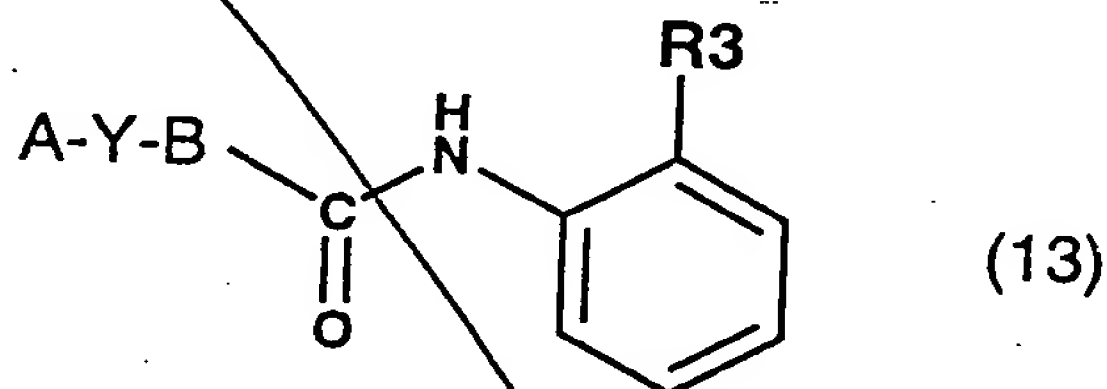
26. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 1 represented by formula (11).



27. A benzamide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 1 represented by formula (12).



28. An anilide having the structure represented by formula (13):



[wherein A and B are independently an optionally substituted phenyl or heterocyclic group which has 1 to 4 substituents selected from the group consisting of a halogen atom, a hydroxyl group, an amino group, a nitro group, a cyano group, an alkyl group having 1 to 4 carbons, an alkoxy group having 1 to 4 carbons, an aminoalkyl group having 1 to 4 carbons, an alkylamino group having 1 to 4 carbons, an acyl group having 1 to 4 carbons, an

acylamino group having 1 to 4 carbons, an alkylthio group having 1 to 4 carbons, a perfluoroalkyl group having 1 to 4 carbons, a perfluoroalkyloxy group having 1 to 4 carbons, a carboxyl group, an alkoxycarbonyl group having 1 to 4 carbons, a phenyl group and a heterocyclic group;

Y is a moiety having -CO-, -CS-, -SO- or -SO₂- which is linear, cyclic or their combination and links A and B;

R³ is a hydroxy or amino group;

the distances between the centroid of ring B (W1), the centroid of ring A (W2) and oxygen or sulfur atom as a hydrogen bond acceptor in the moiety Y (W3) are as follows; W1-W2=6.0 to 11.0 Å, W1-W3=3.0 to 8.0 Å, and W2-W3=3.0 to 8.0 Å.]

or a pharmaceutically acceptable salt thereof.

29. An anilide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 28, wherein A is an optionally substituted heterocycle; R³ is an amino group; and Y is a moiety having -CO- which is linear, cyclic or their combination and links A and B.

30. An anilide derivative or a pharmaceutically acceptable salt thereof as claimed in Claim 29, wherein B is an optionally substituted phenyl; W1-W2 is 7.0 to 9.5 Å; W1-W3 is 3.0 to 5.0 Å; and W2-W3 is 5.0 to 8.0 Å.

31. An anticancer drug comprising one or more compounds as claimed in any of Claims 1 to 30 as active

ingredients.

32. A pharmaceutical composition comprising one or more compounds as claimed in any of Claims 1 to 30 as active ingredients.

suba

add
B1

200607051600